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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/772,253	02/06/2004	Mitsushi Fujiki	042068	6491
38834	7590	08/28/2007	EXAMINER	
WESTERMAN, HATTORI, DANIELS & ADRIAN, LLP			PHAM, THANH V	
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary	Application No.	Applicant(s)
	10/772,253	FUJIKI, MITSUSHI
	Examiner	Art Unit
	Thanh V. Pham	2823

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

1) Responsive to communication(s) filed on 21 August 2007.

2a) This action is **FINAL**. 2b) This action is non-final.

3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

4) Claim(s) 1,2 and 4-10 is/are pending in the application.
4a) Of the above claim(s) _____ is/are withdrawn from consideration.

5) Claim(s) _____ is/are allowed.

6) Claim(s) 1-2 and 4-10 is/are rejected.

7) Claim(s) _____ is/are objected to.

8) Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

9) The specification is objected to by the Examiner.

10) The drawing(s) filed on _____ is/are: a) accepted or b) objected to by the Examiner.

Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).

Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).

11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
a) All b) Some * c) None of:
1. Certified copies of the priority documents have been received.
2. Certified copies of the priority documents have been received in Application No. _____.
3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

1) Notice of References Cited (PTO-892)
2) Notice of Draftsperson's Patent Drawing Review (PTO-948)
3) Information Disclosure Statement(s) (PTO/SB/08)
Paper No(s)/Mail Date _____.
4) Interview Summary (PTO-413)
Paper No(s)/Mail Date _____.
5) Notice of Informal Patent Application
6) Other: _____.

DETAILED ACTION

Continued Examination Under 37 CFR 1.114

1. A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 08/21/2007 has been entered.

Response to Amendment

Claim Rejections - 35 USC § 103

2. The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

3. Claims 1-2 and 5-8 are rejected under 35 U.S.C. 103(a) as being unpatentable over Corvasce et al. US 6,300,654 B1 in combination with Sasaki et al. US 6,444,099 B1 and Matsuura et al. US 6,964,873 B2.

Re claim 1, the Corvasce et al. reference discloses a method of manufacturing a semiconductor device of prior art, comprising:

forming an insulating film 24 over a semiconductor substrate 11;

forming a Ti lower layer 26 of a lower-electrode conductive film on the insulating film 24;

forming an upper layer 7 of the lower-electrode conductive film on the lower layer 26, and constituting a lower-electrode conductive film by the upper and lower layers;

forming a ferroelectric film 17 of PZT or SBT (re claim 7) on the lower-electrode conductive film 7/26;

forming an upper-electrode conductive film 8 on the ferroelectric film 17; and forming a ferroelectric capacitor by patterning the upper-electrode conductive film, the ferroelectric film, and the lower-electrode conductive film, fig. 3.

Re claim 5, the Corvasce et al. reference also discloses the upper layer of the lower-electrode conductive film is a single-layer film made of platinum, col. 3, line 64.

The Corvasce et al. reference does not disclose what method in what temperature used in the process step of forming lower layer of lower-electrode conductive film. In other words, the Corvasce et al. reference does not disclose keeping substrate temperature higher than room temperature and lower than 200 $^{\circ}\text{C}$ while sputtering a Ti lower layer 26 of a lower-electrode conductive film on the insulating film 24.

The Matsuura et al. reference discloses, col. 7 lines 34-50

Referring to FIG. 3A, a SiO₂ film 32 is formed on a Si substrate 31 by a thermal oxidation process with a thickness of 200 nm, for example, and a lower electrode 33 of Pt is formed on the SiO₂ film 32 by a D.C. sputtering process conducted at a room temperature, with an adhesion layer 33A of Ti interposed between the SiO₂ film 32 and the lower electrode 33.

More specifically, (re claim 2) the Ti adhesion layer 33A is formed in an Ar atmosphere under the pressure of 0.7 Pa with a thickness of about 20 nm as represented in TABLE I below. Further, the lower electrode 33 of Pt is formed under the same condition (re claim 5) with a thickness of about 175 nm. The deposition of the Ti film 33A is conducted by setting the D.C. plasma power to 2.6 kW, wherein the deposition of the Ti film 33A is conducted for the duration of 9 seconds while the deposition of the lower electrode 33 is conducted for the duration of 96 seconds while setting the D.C. plasma power to 1.0 kW.

The Sasaki et al. reference discloses, col. 7, lines 15-30

EXAMPLE 1

Sputtering can be carried out under the following conditions as a practical example (hereinafter referred to as the first practical example) of producing a titanium thin film for use as a barrier film. This example pertains to the embodiment given above.

Sputtering power source 3: 13.56 MHz, 8 kW output

Material of target 2: titanium

Type of process gas: argon

Flux of process gas: 120 cc/min

Pressure during film deposition: 60 mTorr

Substrate-biasing voltage: -600 V

Temperature of substrate holder 5 during film deposition: 300.degree. C.

Deposition rate: 500 angstroms/min

Further, The Sasaki et al. reference discloses, col. 4, lines 64-65 "heater 52

controls the temperature of the substrate 50 over a range from room temperature to about 500 °C among many variable parameters of pressure, target size, distance between target and substrate holder, frequency/current of the output of the power source, ratio and flow rates of gases, flux adjusting, electric field, bias voltage, etc. while performing the sputtering process (col's. 3-5).

Choice of temperature, amongst many variable parameters would have been a matter of routine optimization because temperature is known to affect device properties and would depend on the desired device density on the finished wafer and the desired device characteristics. One of ordinary skill in the art would have been led to the recited temperature through routine experimentation to achieve desired deposition and reaction rates.

The Marsuura et al. reference discloses further, (col. 3 lines 38-52)

In general, it is known that the ferroelectric properties of a PZT or PLZT film is related to the orientation of the PZT or PLZT crystals constituting the film.

Commonly, a predominantly (111) or (100)-orientation is obtained for a PZT or PLZT film formed on a Pt lower electrode, which has a self-textured (111)-orientation (*re claim 6*), due to the epitaxial effect, in which the surface energy is minimized as a result of the foregoing film orientation. It should be noted that a PZT or PLZT film has a self-textured (100)-orientation. In order to maximize the remnant polarization of the PZT or PLZT film, it is desired to align the PZT or PLZT crystals, which belong to the tetragonal crystal system, such that the switching direction for the preferential (100)-orientation is perpendicular to the switching electric field.

Meanwhile, it is known that the PZT or PLZT film constituting the ferroelectric capacitor insulation film 16 of FIG. 1 shows a columnar microstructure and that the value of the spontaneous polarization $2 P_r$ is maximized when the crystal grains therein are oriented in the (111) direction.

Re claims 6-8, Matsuura et al.'s PLZT film is formed as the ferroelectric film "by sputtering process contains characteristically low concentration C (carbon)", (col. 8, lines 61-63, *re claim 7*); an orientation direction of the ferroelectric film 34 is a (111) direction, (col. 3, line 39 – col. 4, line 40 and col. 12, lines 10-11, *re claim 8*); and "a Pt lower electrode, which has a self-textured (111)-orientation", col. 3, lines 42-43, [the same as instant Background of the Invention, page 2, "in general, a Pt film oriented in the (222) direction, which is the same direction as the (111) direction, is employed as the lower electrode"], (*re claim 6*).

4. Claims 4 and 10 are rejected under 35 U.S.C. 103(a) as being unpatentable over the combination of Corvasce et al. with Sasaki et al. and Matsuura et al. as applied to claims 1-3 and 5-8 above, and further in view of Ohwaki et al., "Preferred Orientation in Ti Film Sputter-Deposited on SiO_2 Glass: The Role of Water Chemisorption on the Substrate", Jpn. J. Appl. Phys., Vol. 36 (1997) pp L154-L157 (provided by applicant).

The combination teaches substantially all of the instant invention but does not teach crystal orientation, H_2O added during sputtering, ...

The Ohwaki et al. reference discloses a sputtering method (*re claim 2*) for forming Ti (*re claim 3*) on glass which improves the orientation of the Ti film in the preferred (002) direction (*re claim 4*) wherein an orientation with an amount of H₂O (*re claim 10*) to enhance the Ti (002) preferred orientation providing the temperature at 350 °C.

Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to provide the method of Corvasce et al.'s prior art with the conditions of Ohwaki et al. and/or Marsuura et al. because the conditions of Ohwaki et al. and/or Marsuura et al. would provide the ferroelectric capacitor of Corvasce et al. with the Ti (002) preferred orientation for the reliability of the electrode (Ohwaki et al.'s) and with better adhesion (Marsuura et al.'s).

5. Claim 9 is rejected under 35 U.S.C. 103(a) as being unpatentable over the combination of Corvasce et al. with Sasaki et al. and Matsuura et al. as applied to claims 1-3 and 5-8 above, and further in view of Noguchi et al. US 6,716,749 B2.

The combination does not disclose the improvement of the insulating film before forming further the device. The Noguchi et al. reference discloses in col. 21, lines 10-13, quality of the insulating film is improved by exposed a surface of the insulating film to NH₃ plasma. It would have been obvious to one of ordinary skill in the art at the time of the invention to provide the process of the combination with NH₃ plasma nitridation before the lower layer of the lower-electrode conductive film is formed because the plasma nitridation would improve the surface of the insulating film as taught by Noguchi et al.

Response to Arguments

6. Applicant's arguments filed 08/03/2007 have been fully considered but they are not persuasive. The Sasaki et al. reference not only discloses forming titanium nitride film as in example 2 but also forming titanium film in example 1. The disclosures of both Sasaki et al. and Matsuura et al. are thin film of titanium formed in/with different parameters amongst many variable parameters as can be seen in the quotations from the two references. The Matsuura et al.'s discloses "the Ti adhesion layer 33A is formed in an Ar atmosphere under the pressure of 0.7 Pa with a thickness of about 20 nm ... is conducted by setting the D.C. plasma power to 2.6 kW...for the duration of 9 seconds" "at a room temperature" and the Sasaki et al.'s discloses "producing a titanium thin film for use as a barrier film. This example pertains to the embodiment given above." In the condition of "Sputtering power source 3: 13.56 MHz, 8 kW output, Material of target 2: titanium, Type of process gas: argon, Flux of process gas: 120 cc/min, Pressure during film deposition: 60 mTorr, Substrate-biasing voltage: -600 V, Temperature of substrate holder 5 during film deposition: 300.degree. C., Deposition rate: 500 angstroms/min". One of ordinary skill in the art would recognize some parameters are the same and some are different. Thus, One of ordinary skill in the art would have been led to the recited temperature through routine experimentation to achieve desired deposition and reaction rates.

Conclusion

7. The prior art made of record and not relied upon is considered pertinent to applicant's disclosure.

8. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Thanh V. Pham whose telephone number is 571-272-1866. The examiner can normally be reached on M-T (6:30-5:00).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Matthew Smith can be reached on 571-272-1907. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

08/27/2007



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TC 2800
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